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THE COMMISSIONER OF PATENTS AND TRADEMARKS  
Washington, D.C. 20231

Sir:

Transmitted herewith for filing is the patent application of:

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DAISUKE IMANARI, TAKASHI KITAHAMA

Title: MULTI-LAYER EXPANSION-MOLDED ARTICLE OF POLYPROPYLENE RESIN, PRODUCTION PROCESS  
THEREOF, AND CONTAINER, SHOCK-ABSORBING MATERIAL FOR AUTOMOBILE AND AUTOMOTIVE  
MEMBER FORMED OF THE MULTI-LAYER EXPANSION-MOLDED ARTICLE OF POLYPROPYLENE RESIN

Enclosed are:

- ☒ 7 sheets of drawings  
☒ Executed Declaration/Power of Attorney  
☐ Unexecuted Declaration/Power of Attorney  
☒ An Assignment of the invention to: JSP Corporation, Tokyo, Japan

- ☐ Certified copy of:  
☐ Verified statement to establish small entity status  
☐ Preliminary Amendment  
☐ Form PTO-1449 ☐ Information Statement

The filing fee has been calculated as shown below:

			SMALL ENTITY		OTHER THAN A SMALL ENTITY
FOR	NO. FILED	NO. EXTRA	RATE FEE		RATE FEE
BASIC FEE			\$345		\$690
Total Claims	26 -20=	6	x9 =	or	x18= 108
Indep claims	4 -3=	1	x39=	or	x78= 78
<input checked="" type="checkbox"/> Multiple	Dep claims		+130=	or	+260= 260
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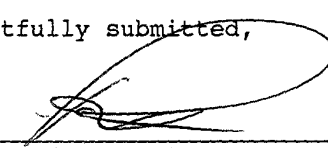
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- ☒ The Commissioner is hereby authorized to charge payment of any additional filing or application processing fees or credit any overpayment to Deposit Account No. 19-1980. Issue and Maintenance fees are excluded from this authorization. A duplicate of this sheet is enclosed.

- ☐ This application is a continuation-in-part of copending application Serial No. \_\_\_\_\_, filed \_\_\_\_\_ and priority is claimed therefrom.

- ☒ Priority is claimed under 35 USC 119 based on:  
Japanese Application 225346 filed August 9, 1999.

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TITLE OF THE INVENTION:

MULTI-LAYER EXPANSION-MOLDED ARTICLE OF  
POLYPROPYLENE RESIN, PRODUCTION PROCESS THEREOF,  
AND CONTAINER, SHOCK-ABSORBING MATERIAL FOR  
5 AUTOMOBILE AND AUTOMOTIVE MEMBER FORMED OF THE  
MULTI-LAYER EXPANSION-MOLDED ARTICLE OF  
POLYPROPYLENE RESIN

## BACKGROUND OF THE INVENTION

10 Field of the Invention:

The present invention relates to a multi-layer expansion-molded article of a polypropylene resin, which has a polypropylene resin layer on the surface of a foamed polypropylene resin layer, and is used as, for example, a light-weight heat insulating panel utilized for a floor, door or the like, a pallet, a container, an automotive member, or the like.

Description of the Background Art:

Multi-layer resinous expansion-molded articles in various forms, in which a thermoplastic resin layer is formed on the surface of a foamed resin layer, have heretofore been known. In recent years, the problem of refuse has been causing great concern, and so there has been a demand for changing containers for fishery products made of an expansion-molded article of polystyrene resin particles, and the like, which have heretofore been disposable, to containers made of a

multi-layer resinous expansion-molded article, in which a polypropylene resin layer is formed on the surface of a foamed polypropylene resin layer, to circulate them as returnable containers capable of sterilizing with steam.

5 Automotive members and the like have also been required to constitute with members made of the same multi-layer resinous expansion-molded article having a polypropylene resin layer on its surface as described above from the viewpoints of recycling ability, strength, lightweight  
10 property, heat insulating property and sound absorbing property.

As this kind of multi-layer resinous expansion-molded articles, there have been known those obtained by a process in which a molded article of a hollow  
15 structure is formed with a thermoplastic resin to use it as a resin layer, and a polyurethane resin is poured into the cavity within the molded article, or foamed beads of a resin are filled thereinto to mold (Japanese Patent Publication No. 10217/1983, Japanese Patent  
20 Application Laid-Open No. 339979/1994, etc.). However, this process has involved problems that the molding process of the multi-layer resinous expansion-molded article becomes complicated, and a special molding machine is required, since a step of forming the resin  
25 layer which becomes a skin and a step of forming a foamed layer in the inside of the resin layer are separately conducted. It has hence been difficult to

provide the multi-layer resinous expansion-molded article on the cheap.

There has also been proposed a process in which a hollow multi-layer parison with a thermoplastic resin layer provided on the surface of a foamed resin layer is molded while compressing it in a mold to cause the opposed inner surfaces of the parison to be fusion-bonded to each other, thereby obtaining a multi-layer resinous expansion-molded article (Japanese Patent Publication No. 27978/1987, Japanese Patent Application Laid-Open No. 312449/1994, etc.).

However, such a process of molding the multi-layer parison while compressing it in the mold as described above has involved a problem that it is difficult to provide any expansion-molded article having a resin layer having a uniform thickness, and so irregularities occur in the surface of the resulting expansion-molded article to impair its appearance, or the resin layer cracks or bores when a molded article having a relatively high draw ratio, such as a container, is formed. In addition, this process has also involved a problem that when a polypropylene resin is used as a base resin, it is difficult to obtain a molded article having a low-density foamed resin layer the density of which is as low as at most  $0.4 \text{ g/cm}^3$ , or when it is attempted to make the density of the foamed resin layer low, the resulting foamed resin layer tends to cause

cell collapse and formation of open cells. As described above, this process has involved some problems to be solved.

## 5 SUMMARY OF THE INVENTION

The present invention has been completed in view of the above-described problems involved in the prior art, and has as its object the provision of a multi-layer expansion-molded article of a polypropylene resin,  
10 which can be produced without need of such complicated processes that a molded article, which will become a resin layer on the surface of a foamed resin layer, is formed, and a foam is then poured thereinto, and is beautiful in appearance and excellent in lightweight  
15 property, mechanical strength, heat insulating property, cushioning property, chemical resistance, recycling ability, etc., a production process thereof and a container, automotive members produced thereby.

According to the present invention, there is thus  
20 provided a multi-layer expansion-molded article of a polypropylene resin, which is obtained by molding a multi-layer parison comprising a foamed resin layer and a resin layer provided on the outer side of the foamed resin layer in a mold in such manner that at least part  
25 of the opposed inner surfaces of the foamed resin layer in the parison are fusion-bonded to each other, and has a polypropylene resin layer on the surface of a foamed

polypropylene resin layer, wherein a melt tension, MT (gf) and a melt flow rate, MFR (g/10 min) obtained by measurement to a polypropylene resin forming the foamed resin layer in the expansion-molded article satisfy the following relationship (1), and a melt tension, MT (gf) and a melt flow rate, MFR (g/10 min) obtained by measurement to a polypropylene resin forming the resin layer on the surface of the foamed resin layer satisfy the following relationship (2) when the melt flow rate, MFR is at least 0.3 (g/10 min), or the melt tension, MT is at least 10 (gf) when the melt flow rate, MFR is not lower than 0.2 (g/10 min), but lower than 0.3 (g/10 min):

$$\log MT > -0.74 \log MFR + 0.66 \quad (1)$$

$$\log MT > -1.02 \log MFR + 0.47 \quad (2)$$

According to the present invention, there is also provided a multi-layer expansion-molded article of a polypropylene resin, which is obtained by molding a multi-layer parison comprising a foamed resin layer and a resin layer provided on the outer side of the foamed resin layer in a mold in such manner that at least part of the opposed inner surfaces of the foamed resin layer in the parison are fusion-bonded to each other, and has a polypropylene resin layer on the surface of a foamed polypropylene resin layer, wherein a melt tension, MT (gf) and a melt flow rate, MFR (g/10 min) obtained by measurement to a polypropylene resin forming the foamed

resin layer in the expansion-molded article satisfy the following relationship (3), and a melt tension, MT (gf) and a melt flow rate, MFR (g/10 min) obtained by measurement to a polypropylene resin forming the resin layer on the surface of the foamed resin layer satisfy the following relationship (4) when the melt flow rate, MFR is at least 0.5 (g/10 min), or the melt tension, MT is at least 10 (gf) when the melt flow rate, MFR is not lower than 0.2 (g/10 min), but lower than 0.5 (g/10 min):

$$\log MT > -0.74 \log MFR + 0.79 \quad (3)$$

$$\log MT > -1.02 \log MFR + 0.69 \quad (4)$$

The expansion-molded article according to the present invention may preferably include a molded article in which the thickness of the propylene resin layer is 100  $\mu\text{m}$  to 10 mm, and the overall density of the expansion-molded article is 20 to 400  $\text{kg/m}^3$ . In the expansion-molded article according to the present invention, the area ratio of the fusion-bonded portion in the inner surface of the expansion-molded article may preferably be at least 25%, more preferably at least 60%, still more preferably at least 80%, particularly preferably at least 95%. The expansion-molded article according to the present invention may further have a skin layer formed of a synthetic resin on the outer side of the resin layer.

According to the present invention, there is

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$$\log \text{MT} > -1.02 \log \text{MFR} + 0.47 \quad (2)$$

According to the present invention, there is still further provided a process for producing a multi-layer



expansion-molded article of a polypropylene resin, which comprises co-extruding a foamable molten resin obtained by adding a foaming agent to a polypropylene resin whose melt tension, MT (gf) and melt flow rate, MFR (g/10 min) satisfy the following relationship (5), and melting and kneading the resultant mixture, and a molten resin obtained by melting and kneading a polypropylene resin whose melt flow rate, MFR (g/10 min) is at least 0.8 (g/10 min), and whose melt tension, MT (gf) and melt flow rate, MFR (g/10 min) satisfy the following relationship (6), or whose melt flow rate, MFR is not lower than 0.2 (g/10 min), but lower than 0.8 (g/10 min) and whose melt tension, MT is at least 10 (gf), thereby forming a multi-layer parison having a resin layer composed of the molten resin on the surface of a foamed resin layer obtained by expanding the foamable molten resin, and then molding the multi-layer parison in a mold in such manner that at least part of the opposed inner surfaces of the foamed resin layer in the parison are fusion-bonded to each other to obtain an expansion-molded article having a polypropylene resin layer on the surface of a foamed polypropylene resin layer:

$$\log MT > -0.74 \log MFR + 1.14 \quad (5)$$

$$\log MT > -1.02 \log MFR + 0.90 \quad (6)$$

In the processes according to the present invention, the polypropylene resin used for the foamable molten resin may preferably have an MFR of 0.3 to 20 g/10 min.

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According to the present invention, there is yet still further provided a shock-absorbing material for automobile formed of the multi-layer expansion-molded article of the polypropylene resin described above, wherein the overall density of the shock-absorbing material is 25 to 300 kg/m<sup>3</sup>, and the thickness of the polypropylene resin layer is 200 μm to 7 mm.

The member for automobile may be a member selected from among a bumper, pillar, instrument panel, spoiler, fender, side step, door trim, grille guard and trunk board.

The above and other objects, features and  
25 advantages of the present invention will become apparent  
from the following description and the appended claims,  
taken in conjunction with the accompanying drawings, in



5 Figs. 1A and 1B. As illustrated in Figs. 1A and 1B, the expansion-molded articles 1 according to the present invention have a structure that a polypropylene resin layer 13 is formed on the surface of an foamed polypropylene resin layer 12.

10           The expansion-molded article 1 according to the present invention is obtained by holding a multi-layer parison in a mold to compression-mold it. The term "parison" as used in the present invention means a tubular or any other hollow plastic formed or molded  
15 article obtained by bonding the opposed ends of two sheets to each other, or extrusion, injection molding or the like and such that generally used in molding such as blow molding. Figs. 2A and 2B illustrate examples of the multi-layer parison 6. The expansion-molded articles 1  
20 illustrated in Figs. 1A and 1B are obtained from the multi-layer parison illustrated in Fig. 2A and the multi-layer parison illustrated in Fig. 2B, respectively.

The multi-layer parison illustrated in Fig. 2A has a structure that a polypropylene resin layer 3 is laminated on the outer surface of a foamed polypropylene resin layer 2, and the multi-layer parison illustrated in Fig. 2B has a structure that a skin layer 4 is

further laminated on the surface of the resin layer 3 of the multi-layer parison shown in Fig. 2A.

As the resin layer 3, is preferably used a non-foamed body, a foamed body having a density of at least 0.4 g/cm<sup>3</sup> or a inorganic material-filled non-foamed body. The multi-layer parison 6 can be obtained by uniting and laminating a foamable molten resin prepared by adding a foaming agent to a polypropylene resin forming a foamed resin layer 2, and melting and kneading the resultant mixture in an extruder, and a molten resin prepared by melting a polypropylene resin forming a resin layer 3 in another extruder (or in order to obtain a multi-layer parison further having a skin layer 4 on the face side of the resin layer 3, a molten resin prepared by melting a resin forming the skin layer 4 in a further extruder) in a die for co-extrusion, and co-extruding them through the die to a lower-pressure zone, thereby expanding the foamable molten resin.

The polypropylene resin layer 13 (hereinafter may be referred to as "the resin layer 13 of the expansion-molded article" or simply "the resin layer 13") in the expansion-molded article 1 according to the present invention corresponds to the resin layer 3 of the multi-layer parison 6. On the other hand, the foamed polypropylene resin layer 12 (hereinafter may be referred to as "the foamed resin layer 12 of the expansion-molded article" or simply "the foamed resin

layer 12") in the expansion-molded article 1 corresponds to the foamed resin layer 2 of the multi-layer parison 6. The skin layer 4 in the multi-layer parison illustrated in Fig. 2B corresponds to the skin layer 14 (hereinafter may be referred to as "the skin layer 14 of the expansion-molded article" or simply "the skin layer 14") provided on the outer side of the resin layer 13 of the expansion-molded article 1 illustrated in Fig. 1B.

As the polypropylene resin used for forming the foamed resin layer 12 of the expansion-molded article, i.e., the polypropylene resin used for forming the foamed resin layer 2 of the multi-layer parison 6, is used a polypropylene resin (hereinafter referred to as "Specific Polypropylene Resin A" simply) comprising a polypropylene resin as a main component, whose melt tension, MT (gf) and melt flow rate, MFR (g/10 min) satisfy the following relationship (1):

$$\log MT > -0.74 \log MFR + 0.66 \quad (1)$$

The use of a polypropylene resin which does not satisfy the relationship (1) fails to form a formed resin layer 2 having a high expansion ratio and a great thickness. Specific Polypropylene Resin A used for forming the foamed resin layer 2 is preferably such that the MT (gf) and MFR (g/10 min) satisfy the following relationship (5). Such a polypropylene resin permits the formation of a foamed resin layer 2 having a higher expansion ratio and a greater thickness.

$$\log MT > -0.74 \log MFR + 1.14 \quad (5)$$

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The MFR of Specific Polypropylene Resin A is preferably 0.3 to 20 g/10 min. If MFR is lower than 0.3 g/10 min, the foam extrusion temperature of a foamable molten resin obtained by melting and kneading such a Specific Polypropylene Resin A and a foaming agent must be made high. As a result, there is a possibility that the viscoelasticity of the resin may be lowered, so that the foaming agent may escape from the resin, resulting in a failure to form a foamed resin layer 2 of a closed cell structure. On the other hand, if MFR exceeds 20 g/10 min, the resultant foamed resin layer 2 of the multi-layer parison undergoes great drawdown due to its high flowability upon heating and molding, and so the provision of a large-sized expansion-molded article requires to take measures against a production apparatus therefor.

As the polypropylene resin used for forming the resin layer 13 of the expansion-molded article, i.e., the polypropylene resin used for forming the resin layer 3 of the multi-layer parison 6, is used a polypropylene resin (hereinafter referred to as "Specific Polypropylene Resin B" simply) comprising a polypropylene resin as a main component, whose melt tension, MT (gf) and melt flow rate, MFR (g/10 min) satisfy the following relationship (2) when the MFR is at least 0.3 (g/10 min) or whose MT is at least 10 (gf)

when the MFR is not lower than 0.1 (g/10 min), but lower than 0.3 (g/10 min):

$$\log MT > -1.02 \log MFR + 0.47 \quad (2)$$

When a polypropylene resin which does not satisfy the relationship (2) when the MFR is at least 0.3 (g/10 min), or a polypropylene resin whose MFR is not lower than 0.1 (g/10 min), but lower than 0.3 (g/10 min) and whose MT is lower than 10 (gf) is used to form a resin layer 3 of the multi-layer parison, the resultant resin layer 3 can not follow the extension of the foamed resin layer 2 upon the molding of such a parison by a molding machine, so that uneven extension occurs in the resin layer 3 to cause a scale pattern, cracks and/or holes in the surface of the resin layer 3. As a result, the resin layer 13 of the resulting expansion-molded article 1 also becomes insufficient. Besides, since any resin having an MFR of lower than 0.1 (g/10 min) is low in flowability, the use of such a low-MFR resin makes it difficult to extrude the molten resin for forming the resin layer 3 unless the extrusion temperature in the die is made high. On the other hand, the foamable molten resin for forming the foamed resin layer 2 requires to lower the foam extrusion temperature in the die until a viscosity suitable for expansion is achieved. Therefore, when the resin having an MFR of lower than 0.1 (g/10 min) is used as a resin for forming the resin layer 3, it is difficult to provide a good multi-layer parison by



co-extruding it with the foamable molten resin for forming the foamed resin layer 2, and the heat of the molten resin for forming the resin layer 3 causes such problems that the cell structure of the foamed resin layer 2 is destroyed. When the MT of Specific Polypropylene Resin B is at least 10 (gf), the poor flowability of Specific Polypropylene Resin B can be offset by its strong tension upon melting so far as the MFR is at least 0.1 (g/10 min). Accordingly, a good expansion-molded article 1 can be provided without causing a scale pattern, cracks, holes and/or the like in the resin layer 13 of the molded article by taking the production conditions of the multi-layer parison, the molding conditions of the parison and the like into adequate consideration. Incidentally, the MFR of Specific Polypropylene Resin B is preferably at most 20 (g/10 min) for the purpose of preventing the drawdown of the resulting multi-layer parison 6. A polypropylene resin whose MFR is at least 0.5 (g/10 min), and whose MT (gf) and MFR (g/10 min) satisfy the following relationship (4), or whose MFR is not lower than 0.2 (g/10 min), but lower than 0.5 (g/10 min), and whose MT is at least 10 (gf) is preferred as Specific Polypropylene Resin B. A polypropylene resin whose MFR is at least 0.8 (g/10 min), and whose MT (gf) and MFR (g/10 min) satisfy the following relationship (6), or whose MFR is not lower than 0.2 (g/10 min), but lower

than 0.8 (g/10 min), and whose MT is at least 10 (gf) is more preferred as Specific Polypropylene Resin B. The use of such a resin B provides a good expansion- molded article 1 having a resin layer 13 evenner in thickness.

$$5 \quad \log MT > -1.02 \log MFR + 0.69 \quad (4)$$

$$\log MT > -1.02 \log MFR + 0.90 \quad (6)$$

The above-described specific resins A and B can be obtained in accordance with, for example, the following process. Namely, such a resin can be obtained by a

10 process in which an ordinary polypropylene resin (the ordinary polypropylene resin used for obtaining Specific Polypropylene Resins A and B will hereinafter be referred to as "the raw polypropylene resin" in order to distinguish it from Specific Polypropylene Resins A and

15 B) low in MT, a peroxide lower in 1-minute half-life temperature than the melting point of the raw polypropylene resin, and a main chain scission-preventing agent are poured into an aqueous medium to stir them, thereby impregnating the raw polypropylene

20 resin with the peroxide and main chain scission-preventing agent such temperature and time conditions that the decomposition of the peroxide is prevented as much as possible, and at least a half of the whole peroxide remains, and the raw resin is then held for a

25 prescribed period of time under temperature conditions that the 1-minute half-life temperature of the peroxide used is reached to decompose to the peroxide, thereby

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slightly crosslinking the raw polypropylene resin in such a manner that its gel content is lower than 1%, or a process in which a low-temperature decomposing peroxide (1-minute half-life temperature: room temperature to about 120°C) is mixed with a raw polypropylene resin containing an atactic component and/or a component that is isotactic, but is not crystalline to heat the resultant mixture at 120°C or lower, thereby bonding the atactic component and/or the non-crystalline isotactic component as a branched chain to the main chain of the raw polypropylene resin to provide a polypropylene resin having long-chain branches. Incidentally, the 1-minute half-life temperature of the peroxide means a temperature at which the half-life of the peroxide becomes 1 minute.

In the above-described process, as the raw polypropylene resin, may be used a propylene homopolymer or a copolymer of propylene with other monomers. Examples of the other monomers include ethylene, 1-butene, isobutylene, 1-pentene, 3-methyl-1-butene, 1-hexene, 3,4-dimethyl-1-butene, 1-heptene and 3-methyl-1-hexene. The copolymer may be either a random copolymer or a block copolymer.

When a copolymer is used as the raw polypropylene resin, it is preferred that a proportion of other monomer components in the copolymer be at most 5.0% by weight for the random copolymer or at most 20.0% by

weight for the block copolymer in order that the properties inherent in the polypropylene resin are not impaired.

Such raw polypropylene resins as described above  
5 may be used either singly or in any combination thereof. With the raw polypropylene resin, may be mixed other resin components such as ethylene resins such as high density polyethylene, low density polyethylene, linear low density polyethylene, linear very low density  
10 polyethylene, ethylene-butene copolymers and ethylene-maleic anhydride copolymers, ethylene-vinyl acetate copolymers, and styrene resins within limits not impeding the properties inherent in polypropylene as needed.

15 Examples of the peroxide used for preparing Specific Polypropylene Resins A and B from the raw polypropylene resin include lauroyl peroxide, m-toluoyl-benzoyl peroxide, benzoyl peroxide and bis(4-butylcyclohexyl) peroxydicarbonate. Among these, a peroxide whose  
20 1-minute half-life temperature is lower than the melting point of the raw polypropylene resin used is selected. The main chain scission-preventing agent serves to prevent the main chain of the raw polypropylene resin being cut by the peroxide. For example, methyl  
25 methacrylate, divinylbenzene, triallyl cyanurate or the like is used. As the aqueous medium, is generally water to which a surfactant has been added.

Specific Polypropylene Resin A satisfying the relationship (1) between MT and MFR can be obtained by controlling the length of branched chains and the proportion of branch points of the polypropylene resin having branched chains obtained in the above-described process. In particular, the length of the branched chains is lengthened, and the proportion of the branch points is relatively lessened, thereby obtaining Specific Polypropylene Resin A satisfying the relationship (5). Besides, Specific Polypropylene Resin B satisfying the relationship (2) between MT and MFR can be prepared in the same manner as in the above-described Specific Polypropylene Resin A or may be obtained by improving its flowability with a lubricant or the like even in a polypropylene resin of a linear structure. In particular, a polypropylene resin having branched chains is selected for preparation, thereby obtaining Specific Polypropylene Resin B satisfying the relationship (4) or (6).

However, the above-described processes are examples for obtaining Specific Polypropylene Resins A and B, and Specific Polypropylene Resins A and B respectively used as base resins for forming the foamed resin layer 12 of the expansion-molded article and the resin layer 13 thereof are not limited to those obtained by the above-described processes, and no particular limitation is imposed on the procurement methods thereof



by mixing another polymer must satisfy the relationship between MT and MFR in Specific Polypropylene Resins A and B.

As other polymers capable of mixing with Specific Polypropylene Resins A and B, may be used the same resins as those exemplified as other resin components capable of mixing with the raw polypropylene resin in the above-described processes as examples for obtaining Specific Polypropylene Resins A and B, and besides rubber such as ethylene-propylene rubber; polystyrene elastomers such as styrene-butadiene-styrene block copolymers, styrene-isoprene-styrene block copolymers and hydrogenated products thereof; and polyolefin elastomers comprising a polyolefin copolymer such as a propylene-ethylene block copolymer as a hard segment and polyolefin copolymer rubber such as an ethylene-propylene random copolymer or ethylene-propylene-diene random copolymer as a soft segment; and the like.

Incidentally, when a proportion of other polymer components than the polypropylene component in Specific Polypropylene Resin A or B is increased, the properties inherent in the polypropylene resin may possibly be impaired even when such a polymer satisfies the relationship between MT and MFR. Therefore, when other polymers are mixed with Specific Polypropylene Resin A or B, the mixture is prepared so as to contain the polypropylene component in a proportion of at least 65%

Incidentally, Specific Polypropylene Resins A and B used in the present invention may be those prepared in such a manner that the MT and MFR of the resulting mixtures satisfy the respective relationships between MT and MFR by suitably mixing other polymers than the raw polypropylene resin with the raw polypropylene resin.

The melt tension (MT) in the present specification  
can be measured by means of a Melt Tension Tester II  
Model manufactured by Toyo Seiki Seisaku-Sho, Ltd. or  
the like, which has been produced in accordance with  
ASTM D 1238. Specifically, a melt tension tester  
equipped with a cylindrical orifice having a straight  
hole having a hole diameter of 2.09 mm and a length of 8  
mm is used, a resin sample is extruded into a strand  
through the orifice under conditions of a resin  
temperature of 230°C and a piston speed of 10 mm/min to  
hang the strand on a pulley for detection of tension  
having a diameter of 45 mm, and the strand is taken up  
on a take-up roll having a diameter of 50 mm while  
gradually increasing take-up speed in a proportion of  
about 5 rpm/sec (take-up acceleration of strand: about  
 $1.3 \times 10^{-2}$  m/sec<sup>2</sup>). The take-up speed is first increased  
until the strand hanged on the pulley for detection of  
tension is broken to determine a take-up speed R (rpm)  
at the time the strand has been broken. The strand is



then taken up at a fixed take-up speed of  $R \times 0.7$  (rpm) to measure a melt tension (MT) of the strand detected by a detector connected to the pulley for detection of tension with time and chart it taking MT (gf) on an axis of ordinate and time (second) on an axis of abscissa, thereby obtaining a graph having such amplitude as illustrated in Fig. 3. The MT in the present invention means a median (X) of amplitude at a stable amplitude portion. However, when the strand is not broken even at the time the take-up speed has reached 500 (rpm), a value determined as described above from a graph obtained by take-up at a take-up speed of 500 (rpm) is regarded as a melt tension (MT). Incidentally, specific amplitude occurred rarely on the above graph is ignored. With respect to the melt flow rate (MFR), a value obtained by the measurement at 230°C under a load of 2.16 kgf described in JIS K 7210 is adopted.

In the case of the expansion-molded article 1 further having the skin layer 14 on the outer surface of the resin layer 13, examples of a resin or polymer (hereinafter referred to as resin simply) used for forming the skin layer 14 include thermoplastic resins, such as polyethylene resins such as low density polyethylene, high density polyethylene and linear low density polyethylene; polypropylene resins such as homopolypropylene, propylene-ethylene random copolymers, propylene-1-butene random copolymers, propylene-



incorporated into the foamed resin layer 12, resin layer 13 and skin layer 14 of the expansion-molded article 1 as needed. It is particularly preferable that an inorganic material such as short fiber or clay be subjected to a surface treatment and added to the respective layers, thereby enhancing stiffness of the layers to provide an expansion-molded article 1 having high stiffness.

The thickness of the resin layer 13 of the expansion-molded article 1 according to the present invention is preferably 100  $\mu\text{m}$  to 10 mm, more preferably 100  $\mu\text{m}$  to 7 mm. If the thickness of the resin layer 13 is too small, the molded article 1 has a possibility that the surface smoothness thereof may be deteriorated to make the surface appearance thereof poor. If the thickness of the resin layer 13 is too great, the whole weight of the expansion-molded article 1 becomes heavy, so that it is hard to say that the molded article is light-weight according to the use of the molded article 1, and moreover the closed cell content in the foamed resin layer 12 of the expansion-molded article 1 may lower in some cases. More specifically, since the resin layer and the foamed resin layer are simultaneously extruded through a co-extrusion die by a co-extrusion process, the quantity of heat of a molten resin forming the resin layer becomes high when the thickness of the resin layer is great, so that the foamed resin layer is



A multi-layer parison of a layer structure composed of 3 or more layers like the multi-layer parison illustrated in Fig. 2B used for obtaining an expansion-molded article 1 further having a skin layer 14 on the surface of the resin layer 13 of the expansion-molded article can be obtained by using the same number of extruders as the number of layers and a ring die for co-extrusion of a structure that a plurality of resins can be united and laminated according to the number of layers in the multi-layer parison.

As a foaming agent added to Specific Polypropylene Resin A for forming the foamed resin layer 12 of the expansion-molded article 1, may be used either of a volatile foaming agent or a decomposable foaming agent. In order to obtain a foamed resin layer having a high expansion ratio, however, it is preferable to use a volatile foaming agent. Examples of the volatile foaming agent include aliphatic hydrocarbons such as propane, n-butane, isobutane, n-pentane, isopentane, n-hexane, isohexane and cyclohexane; chlorinated hydrocarbons such

as methyl chloride and ethyl chloride; and fluorinated hydrocarbons such as 1,1,1,2-tetrafluoroethane and 1,1-difluoroethane. Examples of the decomposable foaming agent include azodicarbonamide. The foaming agents  
5 mentioned above may be used in combination of two or more thereof. A decomposable foaming agent also having a function as a cell stabilizer may be used in combination with a volatile foaming agent.

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10 The production process of an expansion-molded article 1 according to the present invention will hereinafter be described in more detail with reference to Figs. 4 and 5. As illustrated in Fig. 4, for example, a foamable molten resin and molten resin(s) for forming a foamed resin layer, a resin layer 3 and a skin layer 4  
15 provided as needed are respectively prepared in separate extruders (not illustrated), and these molten resins are extruded from the extruders to a lower-pressure zone while uniting and laminating them is a die 21, thereby obtaining a multi-layer parison 6. An accumulator may be  
20 provided between the extruders and the die 21 or within the die 11 as needed.

A mold split in a male mold 22a and a female mold 22b is closed so as to hold the multi-layer parison 6 formed by the extrusion through the die 21 from its both  
25 sides. When the mold is closed, the multi-layer parison 6 is compressed on the inner surface of the mold to bring the multi-layer parison 6 into close contact with

When a pipe 23 for pressure reduction is provided in the mold as illustrated in Fig. 4 to conduct the molding while reducing the pressure between the inner surface of the mold and the outer surface of the parison, the outer surface of the multi-layer parison 6 can be fully brought into close contact with the inner surface of the mold to obtain an expansion-molded article 1 conforming to the shape of the mold with good reproducibility and having good appearance. When a mold having a pattern in its inner surface is used, the transferability of the pattern to the resulting expansion-molded article becomes good, and so the

appearance thereof is far excellent.

There is also adopted a process comprising closing an open under end of the multi-layer parison 6 by a parison pinch just after the extrusion of the multi-layer parison 6 through the die 21, blowing air into the parison 6 to inflate it and then conducting molding in the same manner as described above.

Incidentally, Figs. 4 and 5 conceptually illustrate an example of the production process of the expansion-molded article 1 according to the present invention, and the specific procedure for producing the expansion-molded article 1 according to the present invention is not limited to that illustrated. When the multi-layer parison 6 is subjected to deep drawing, a female mold 22b is constructed by a bottom part 24 and side parts 25 provided on 4 sides of the bottom part 24 in such a manner that the set-up and development of the female mold 22b can be conducted by moving the side parts 25. It is preferred that in the course of closing a male mold 22a and the female mold 22b, the molding be conducted while setting up the female mold 22b from a developed state.

In the present specification, a proportion of the fusion-bonded portion of the opposed foamed resin layers in the inner surface of the multi-layer parison 6 is expressed by an area ratio (%) of the fusion-bonded portion. In the present invention, the area ratio of the

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fusion-bonded portion in the expansion-molded article 1 is preferably at least 25%, preferably at least 60%, more preferably at least 80%, particularly preferably at least 95% from the viewpoint of achieving sufficient mechanical strength and dimensional stability. The area ratio of the fusion-bonded portion can be determined in the following manner.

The expansion-molded article 1 is cut at a plane perpendicular to the extrusion direction of the multi-layer parison 6 used for producing the expansion-molded article so as to divide it into 10 equal pieces. The cut pieces of the expansion-molded article are then arranged in order that the expansion-molded article is made up, thereby determining a length (mm) of the fusion-bonded portion of the opposed foamed resin layers in the inner surface of the multi-layer parison at a right sectional surface in each cut piece of the molded article. A value obtained by doubling this length is regarded as a length,  $L$  (mm) of the fusion-bonded portion. On the other hand, when a part of the opposed foamed resin layers in the inner surface of the multi-layer parison is not fusion-bonded to each other to form a void portion, a length,  $M$  (mm) of the inner circumferential edge of the void portion is determined. The length,  $L$  of the fusion-bonded portion and the length,  $M$  of the inner circumferential edge of the void portion are determined as the respective totals of all the fusion-bonded

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found from the foamed resin layer 12 and the resin layer 13 of the expansion-molded article may differ from the values of MT and MFR of the original Specific Polypropylene Resins A and B due to differences in heat history up to the provision of the expansion-molded article 1, and the like. Besides, even in expansion-molded articles obtained by using the same Specific Polypropylene Resins, the values of MT and MFR found from the foamed resin layers 12 and the resin layers 13 of the resultant expansion-molded articles may differ from each other due to differences in extrusion conditions upon the production of the multi-layer parison, and the like. Incidentally, the MT and MFR of the foamed resin layer 12 of the expansion-molded article are determined by using, as a sample for measurement, that obtained by placing a sample cut out of the foamed resin layer for about 15 minutes in a vacuum oven at 200°C to heat and melt it, thereby defoaming it. The values of MT and MFR of the foamed resin layer 12 and the resin layer 13 determined from a sample cut out of the expansion-molded article are values determined in accordance with the same method as in the measurement of MT and MFR of Specific Polypropylene Resins A and B.

In the expansion-molded article 1 according to the present invention, in spite of the heat history in the course of the production of the expansion-molded article

1 according to the present invention, it is required  
 from the viewpoint of achieving the intended object of  
 the present invention that when MT and MFR are  
 determined from samples cut out of the foamed resin  
 5 layer 12 and the resin layer 13 in the above-described  
 manner, the melt tension, MT (gf) and the melt flow rate,  
 MFR (g/10 min) obtained by measurement to the  
 polypropylene resin forming the foamed resin layer 12 in  
 the expansion-molded article satisfy the following  
 10 relationship (1), and moreover the melt flow rate, MFR  
 obtained by measurement to the polypropylene resin  
 forming the resin layer 13 of the expansion-molded  
 article is at least 0.3 (g/10 min), and the melt tension,  
 MT (gf) and the melt flow rate, MFR (g/10 min) obtained  
 15 by measurement to the polypropylene resin forming the  
 resin layer 13 satisfy the following relationship (2),  
 or the melt flow rate, MFR obtained by measurement to  
 the polypropylene resin forming the resin layer 13 of  
 the expansion-molded article is not lower than 0.2 (g/10  
 20 min), but lower than 0.3 (g/10 min), and the melt  
 tension, MT obtained by measurement to the polypropylene  
 resin forming the resin layer 13 is at least 10 (gf):

$$\log MT > -0.74 \log MFR + 0.66 \quad (1)$$

$$\log MT > -1.02 \log MFR + 0.47 \quad (2)$$

25 In order to provide an expansion-molded article  
 excellent in appearance and mechanical properties even  
 in a large-sized expansion-molded article, a deep-draw

5 forming the foamed resin layer 12 in the expansion-  
molded article satisfy the following relationship (3),  
and moreover the melt flow rate, MFR obtained by  
measurement to the polypropylene resin forming the resin  
layer 13 of the expansion-molded article is at least 0.5  
10 (g/10 min), and the melt tension, MT (gf) and the melt  
flow rate, MFR (g/10 min) obtained by measurement to the  
polypropylene resin forming the resin layer 13 satisfy  
the following relationship (4), or the melt flow rate,  
MFR obtained by measurement to the polypropylene resin  
5 forming the resin layer 13 of the expansion-molded  
article is not lower than 0.2 (g/10 min), but lower than  
0.5 (g/10 min), and the melt tension, MT obtained by  
measurement to the polypropylene resin forming the resin  
layer 13 is at least 10 (gf):

$$\log MT > -1.02 \log MFR + 0.69 \quad (4)$$

Since the expansion-molded article according to the present invention is obtained by molding the multi-layer parison 6 having the resin layer 3 composed of a polypropylene resin on the outer surface of the foamed resin layer 2 composed of a polypropylene resin, the adhesion of the foamed resin layer 12 to the resin layer

In the expansion-molded article 1 according to the present invention, the density of the foamed resin layer 12 is preferably 20 to 500 kg/m<sup>3</sup>, more preferably 25 to 400 kg/m<sup>3</sup>. If the density of the foamed resin layer 12 exceeds 500 kg/m<sup>3</sup>, it is hard to say that the molded article is light-weight, and physical properties such as heat insulating property may possibly be deteriorated. If it is attempted to make the density of the foamed resin layer 12 lower than 20 kg/m<sup>3</sup>, the cell membrane making up the foamed resin layer 2 of the multi-layer parison becomes too thin, resulting in a possibility that molding may be difficult due to easy collapse of cells upon molding of the parison, and so the expansion-molded article 1 may not be provided as a good product.

25           The density of the foamed resin layer 12 in the  
expansion-molded article 1 is determined in accordance  
the following method. A portion of the foamed resin

layer 12 is cut out of the expansion-molded article 1, the weight (g) of the sample cut out is divided by a volume (cm<sup>3</sup>) found from the external size of the sample to convert the value found into a kg/m<sup>2</sup> unit. Since the density of the foamed resin layer 12 may vary according to the place measured, samples are cut out of the foamed resin layer 12 of the expansion-molded article at 10 or more places in different portions such as a bottom and sides, and the density of each sample is measured, and the maximum and minimum values are excluded from the resultant measured values to take an arithmetic mean of the remaining values, thereby regarding the average value as the density of the foamed resin layer 12 of the expansion- molded article.

The thickness of the resin layer 13 of the expansion-molded article 1 according to the present invention is preferably 100  $\mu$ m to 10 mm. If the thickness of the resin layer 13 is too small, the molded article 1 has a possibility that the surface smoothness thereof may be deteriorated to make the surface appearance thereof poor. If the thickness of the resin layer 13 is too great on the other hand, the whole weight of the expansion-molded article 1 becomes heavy, so that it is hard to say that the molded article is light-weight according to the use of the molded article 1. The thickness of the resin layer 13 is determined by measurement of the section of the molded article 1. In

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this case, the measurement may also be conducted by enlarging the section through a microscope as needed, and using an enlarged projection drawing. Since the thickness of portions of the resin layer 13 situated at the upper and lower, and right and left peripheries of the expansion-molded article 1 near mold clamps for the parison (X in Fig. 5) may exhibit a small value, and so it is preferred that the thickness of another main portion than those near the clamps should fall within the above range.

The overall density of the expansion-molded article 1 according to the present invention is generally 20 to 800 kg/m<sup>3</sup>, preferably 20 to 400 kg/m<sup>3</sup>. If the overall density of the expansion-molded article 1 exceeds 800 kg/m<sup>3</sup>, the lightweight property and heat insulating property of the molded article may possibly be deteriorated. If the overall density of the molded article 1 is lower than 20 kg/m<sup>2</sup> on the other hand, the mechanical properties such as compressive strength of the molded article may possibly be deteriorated.

The overall density of the expansion-molded article 1 is a value determined by dividing the weight (kg) of the expansion-molded article 1 by its volume (m<sup>3</sup>) measured by sinking the molded article 1 in water. However, even when void portions defined by non-fusion-bonded portions at the inner surface of the multi-layer parison are present in the interior of the molded



article 1, the volume of such void portions is not subtracted from the value determined as the volume of the molded article.

5 Figs. 1A and 1B illustrate sectional shapes of specifically exemplified containers formed of the expansion-molded article 1 according to the present invention, in which the multi-layer parison is molded in such a manner that the opposed innermost foamed layers of the multi-layer parison are completely fusion-bonded to each other without any space, and a container space is defined.

10 In such containers as described above, those in which the overall density is 30 to 400 kg/m<sup>3</sup> are excellent in lightweight property and heat insulating property. Besides, those in which the thickness of the resin layer 13 is 200 to 5 mm are excellent in high durability and surface smoothness, well balanced between appearance and lightweight property, undergo only minimal shrinkage after molding and have excellent mechanical strength against external force. Further, they can be sterilized with steam due to their excellent heat resistance and are hence particularly suitable for use as returnable containers.

25 Fig. 7 is a horizontal sectional view of an automotive bumper illustrated as a specific example of a shock-absorbing material for automobile formed of the expansion-molded article 1 according to the present

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invention. The expansion-molded article 1 illustrated in Fig. 7 is obtained by compression-molding the multi-layer parison 6, and has a structure that the opposed inner-side foamed resin layers of the multi-layer parison 6 are fusion-bonded to each other at the central portion and both ends of the molded article 1 and not fusion-bonded to form void portions between the central portion and both ends. When the expansion-molded article 1 according to the present invention is used as a shock-absorbing material for automobile such as the above-described bumper, the overall density thereof is controlled to 25 to 300 kg/m<sup>3</sup>, thereby providing a shock-absorbing material which is light-weight and excellent in energy-absorbing efficiency. Besides, the thickness of the resin layer 13 is controlled to 200 to 7,000 μm, thereby providing a shock-absorbing material having excellent surface smoothness and impact resistance. The volume of the shock-absorbing material formed of the molded article according to the present invention is preferably 5,000 to 400,000 cm<sup>3</sup>.

The present invention will hereinafter be described in more detail by the following Examples. The melt tensions, MT (gf) and melt flow rates, MFR (g/10 min) of polypropylene resins used in the following Examples and Comparative Examples are shown in Table 1.

Table 1

	Polypropylene resin	MT (gf)	MFR (g/10 min)
Resin A	SD632 (propylene resin); product of Montell Canada Co.	20	3
Resin B	PF814 (homopolymer); product of Montell Canada Co.	22	3
Resin C	PM620A (propylene-ethylene random copolymer); product of Nippon Polyolefin K.K.	0.4	8
Resin D	J700GP (homopolymer); product of Idemitsu Petrochemical Co., Ltd.	0.4	8
Resin E	SB7000 (propylene-ethylene block copolymer); product of Chisso Corporation	26	0.12
Resin F	EG7F (propylene-ethylene random copolymer); product of Nippon Polychem K.K.	1.6	2
Resin G	E150GK (propylene-ethylene block copolymer); product of Idemitsu Petrochemical Co., Ltd.	3.9	0.6
Resin H	Mixture of Resin A/Resin C = 50/50 (weight ratio)	8	6
Resin I	Mixture of Resin B/Resin D = 50/50 (weight ratio)	8	6

## Example 1:

Into 100 parts by weight of Resin A, were  
 5 incorporated 3 parts by weight of a master batch of cell  
 stabilizer prepared by blending 5 parts by weight of  
 sodium citrate and 10 parts by weight of talc into 100  
 parts by weight of a low density polyethylene resin. The  
 resultant mixture was fed to an extruder having a bore

diameter of 65 mm, and isobutane as a volatile foaming agent was forced in a proportion of 2 parts by weight per 100 parts by weight of Resin A into the resin in the extruder and kneaded to prepare a foamable molten resin.

5           On the other hand, a colorant was mixed into Resin A, and the mixture was fed to an extruder having a bore diameter of 40 mm and heated and melted to prepare a non-foamable molten resin. After the molten resins melted and kneaded in the respective extruders were then  
10 charged into separate accumulators connected to the extruders, the foamable molten resin and the non-foamable molten resin were injected from the respective accumulators so as to cause them to join in a die, and to laminate a resin layer formed of the non-foamable  
15 molten resin on the outer surface of a foamed resin layer formed by expansion of the foamable molten resin, and extruded through the die to form a multi-layer parison with the resin layer laminated on the outer surface of the foamed resin layer.

20           A box-shaped split mold arranged right under the die was then closed so as to hold the multi-layer parison thus extruded as illustrated in Figs. 6A and 6B, thereby deforming the multi-layer parison into a flat form by the inner surface of the mold, and the pressure  
25 within the mold was reduced to bring the multi-layer parison into close contact with the inner surface of the mold, thereby molding it. Thereafter, the resultant

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molded article was cooled and released from the mold to obtain an expansion-molded article in the form of a truncated cone-like container having an external size at an opening part of 300 mm x 260 mm, an external size at a bottom part of 260 mm x 180 mm, a height of 110 mm and a wall thickness of about 20 mm. As the mold, was used that subjected to a satinizing processing at its inner surface. The expansion-molded article thus obtained had good surface profile, and a satin pattern on the inner surface of the mold finely appeared on the surface of the molded article. In addition, appearance failure caused by shrinkage deformation, buckling and/or the like was not observed. The section of the molded article was observed. As a result, the molded article was found to have a resin layer formed of the outer resin layer of the multi-layer parison on the surface of the foamed resin layer of the expansion-molded article, and the opposed inner surfaces of the foamed resin layer in the multi-layer parison were fully fusion-bonded to each other without any space.

The foamed resin layer was cut out of the expansion-molded article to measure the MFR and MT of the resin forming the foamed resin layer, thereby finding values of  $\log MT$  and  $-0.74 \log MFR + 0.66$ . The values are shown in Table 2. Further, the resin layer was cut out of the expansion-molded article to measure the MFR and MT of the resin forming the resin layer,

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Table 2

	Resin forming foamed resin layer of expansion-molded article				
	Kind of resin	MFR (g/10 min)	MT (gf)	Value of log (MT)	Value of $-0.74\log(\text{MFR}) + 0.66$
Ex. 1	Resin A	4.5 (3)	6 (20)	0.78 (1.30)	0.18 (0.31)
Ex. 2	Resin A	4.5 (3)	6 (20)	0.78 (1.30)	0.18 (0.31)
Ex. 3	Resin A	4.5 (3)	6 (20)	0.78 (1.30)	0.18 (0.31)
Ex. 4	Resin B	8.1 (3)	1.6 (22)	0.20 (1.34)	0.012 (0.31)
Ex. 5	Resin B	8.1 (3)	1.6 (22)	0.20 (1.34)	0.012 (0.31)
Ex. 6	Resin B	8.1 (3)	1.6 (22)	0.20 (1.34)	0.012 (0.31)
Ex. 7	Resin B	8.1 (3)	1.6 (22)	0.20 (1.34)	0.012 (0.31)
Ex. 8	Resin A	4.5 (3)	6 (20)	0.78 (1.30)	0.18 (0.31)
Ex. 9	Resin A	4.5 (3)	6 (20)	0.78 (1.30)	0.18 (0.31)
Ex. 10	Resin A	4.4 (3)	6.5 (20)	0.81 (1.30)	0.18 (0.31)
Comp. Ex. 1	Resin A	4.5 (3)	6 (20)	0.78 (1.30)	0.18 (0.31)
Comp. Ex. 2	Resin C	8.5 (8)	0.4 (0.4)	-0.40 (-0.40)	0.028 (-0.008)
Comp. Ex. 3	Resin F	2.6 (2)	1.5 (1.6)	0.18 (0.20)	0.35 (0.44)

Table 3

	Resin forming resin layer of expansion-molded article				
	Kind of resin	MFR (g/10 min)	MT (gf)	Value of log(MT)	Value of $-1.02\log(\text{MFR}) + 0.47$
Ex. 1	Resin A	5.6 (3)	5.0 (20)	0.70 (1.30)	-0.29 (-0.017)
Ex. 2	Resin C	8.4 (8)	0.4 (0.4)	-0.40 (-0.40)	-0.47 (-0.45)
Ex. 3	Resin H	7.1 (6)	4.1 (8)	0.61 (0.90)	-0.40 (-0.32)
Ex. 4	Resin B	10.1 (3)	0.4 (22)	-0.40 (1.34)	-0.55 (-0.017)
Ex. 5	Resin I	7.1 (6)	3.5 (8)	0.54 (0.90)	-0.40 (-0.32)
Ex. 6	Resin D	6.4 (8)	6.0 (0.4)	0.78 (-0.40)	-0.35 (-0.45)
Ex. 7	Resin E	0.3 (0.12)	16.4 (26)	1.21 (1.41)	1.00 (1.40)
Ex. 8	Resin F	2.8 (2)	1.6 (1.6)	0.20 (0.20)	-0.013 (0.16)
Ex. 9	Resin H	7.1 (6)	4.1 (8)	0.61 (0.90)	-0.40 (-0.32)
Ex. 10	Resin A	5.5 (3)	5.4 (20)	0.74 (1.30)	-0.29 (-0.017)
Comp. Ex. 1	Resin G	0.9 (0.6)	2.8 (3.9)	0.45 (0.59)	0.52 (0.70)
Comp. Ex. 2	Resin H	7.1 (6)	4.1 (8)	0.61 (0.90)	-0.40 (-0.32)
Comp. Ex. 3	Resin H	7.1 (6)	4.1 (8)	0.61 (0.90)	-0.40 (-0.32)

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Table 4

	Expansion-molded article						
	Thickness of resin layer (mm)		Thickness of *3 foamed resin layer (mm)		Density of foamed resin layer (kg/m <sup>3</sup> )	Overall density of expansion-molded article (kg/m <sup>3</sup> )	Appearance
	Bottom	Side	Bottom	Side			
Ex. 1	1.2	0.9	16	16	150	250	◎
Ex. 2	1.1	0.9	16	16	150	240	◎
Ex. 3	1.2	0.9	16	16	150	250	◎
Ex. 4	1.2	0.9	16	16	130	230	○
Ex. 5	1.1	0.9	16	16	130	220	◎
Ex. 6	1.1	0.9	16	16	130	220	◎
Ex. 7	1.2	0.9	16	16	170	270	○
Ex. 8	1.2	0.9	16	16	150	260	△
Ex. 9	0.9	0.8	19	14	150	240	◎
Ex. 10	35 *4		50 *5		180	210	◎
Comp. Ex. 1	1.2	0.9	16	16	150	260	*1
Comp. Ex. 2	*2	*2	*2	*2	*2	*2	*2
Comp. Ex. 3	*2	*2	*2	*2	*2	*2	*2

- 5

Incidentally, the evaluation of appearance in Table 4 was conducted by visually observing a container-like expansion-molded article sample to judge the appearance in accordance with the following standard:

- 5           ◎: Excellent in surface smoothness and reproducibility to a mold without causing a scale pattern, cracks and/or holes;
- : Causing none of scale pattern, cracks and holes;
- 10          △: Resin melt flow marks are slightly observed on a part of the side wall of the container-like expansion-molded article though neither cracks or holes were observed.

Examples 2 to 6, 8 and 9:

- 15           Multi-layer parisons were separately obtained in the same manner as in Example 1 except that their corresponding resins for forming a foamed resin layer of an expansion-molded article shown in Table 2 and their corresponding resins for forming a resin layer of an
- 20   expansion-molded article shown in Table 3 were respectively used. Each of the parisons was molded to obtain a container-like expansion-molded article. The expansion-molded articles thus obtained had good appearance like Example 1. In the container-like
- 25   expansion-molded articles, the opposed inner surfaces of the foamed resin layer in the multi-layer parison were fully fusion-bonded to each other without any space. The

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various properties of the expansion-molded articles are shown in Tables 2 to 4. In Example 4, a mixed volatile foaming agent composed of 30% by weight of isobutane and 70% by weight of n-butane was used in place of isobutane used as the volatile foaming agent in Example 1.

Example 7:

An expansion-molded article was obtained in the same manner as in Example 1 except that a multi-layer parison obtained in the same manner as in Example 1 except that its corresponding resin for forming a foamed resin layer of an expansion-molded article shown in Table 2 and its corresponding resin for forming a resin layer of an expansion-molded article shown in Table 3 were used, and the amount of the volatile foaming agent was changed to 1.8 parts by weight per 100 parts by weight of Resin B and the expansion-molded article was molded by means of a box-shaped multipart mold different from the mold used in Example 1. The expansion-molded articles thus obtained was not observed causing appearance failure by shrinkage deformation, buckling and/or the like, but had good appearance. In the expansion-molded articles, the opposed inner surfaces of the foamed resin layer in the multi-layer parison were fully fusion-bonded to each other without any space. The various properties of the expansion-molded articles are shown in Tables 2 to 4. In Example 7, the box-shaped multipart mold was used as a mold for molding the multi-

layer parison to form an expansion-molded article in the form of a rectangular-parallelepiped container having an external size at an opening part of 300 mm x 230 mm, a height of 120 mm and a wall thickness of about 20 mm.

5 Example 10:

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A multi-layer parison was formed in the same manner as in Example 1 except that the amount of the foaming agent was changed to 1.9 parts by weight. A bumper-shaped mold arranged right under the die was then closed so as to hold the multi-layer parison thus extruded as illustrated in Figs. 4 and 5, thereby deforming the multi-layer parison into like a flat form by the inner surface of the mold, and the pressure within the mold was reduced to bring the multi-layer parison into close contact with the inner surface of the mold, thereby molding it. Thereafter, the resultant molded article was cooled and released from the mold to obtain a shock-absorbing material for automobile in the form of a bumper having an external size of width = 1,400 mm, height = 260 mm, maximum thickness,  $h_1$  at a hollow part in fig.7 = 110 mm and wall thickness,  $h_2$  portion in fig.7 = about 60 mm. The expansion-molded article thus obtained had good surface profile, and appearance failure caused by shrinkage deformation, buckling and/or the like was not observed. The section of the molded article was observed. As a result, the molded article was found to have a resin layer formed of

the outer resin layer of the multi-layer parison on the surface of the foamed resin layer of the expansion-molded article, and the area ratio of the fusion-bonded portion in the inner surface of the expansion-molded article was 66%.

The foamed resin layer was cut out of the expansion-molded article to measure the MFR and MT of the resin forming the foamed resin layer, thereby finding values of  $\log MT$  and  $-0.74 \log MFR + 0.66$ . The values are shown in Table 2. Further, the resin layer was cut out of the expansion-molded article to measure the MFR and MT of the resin forming the resin layer, thereby finding values of  $\log MT$  and  $-1.02 \log MFR + 0.47$ . The values are shown in Table 3. Further, the thickness of the resin layer, thickness of the foamed resin layer and density of the foamed resin layer of the expansion-molded article, and the overall density of the expansion-molded article were measured, and the results thereof are shown in Table 4. Incidentally, numerical values indicated in parentheses in numerical values shown in Tables 2 and 3 are values in the original resins (base resins) used for producing the respective multi-layer parisons.

#### Comparative Example 1:

A multi-layer parison was obtained in the same manner as in Example 1 except that its corresponding resin for forming a foamed resin layer of an expansion-

molded article shown in Table 2 and its corresponding resin for forming a resin layer of an expansion-molded article shown in Table 3 were used, and the parison thus obtained was molded in the same manner as in Example 1 to obtain a container-like expansion-molded article. The molded article thus obtained was observed causing break at a part of the resin layer and partially exposing the foamed resin layer on the surface. The various properties of the container-like expansion-molded article are shown in Tables 2 to 4.

Comparative Examples 2 and 3:

Their corresponding resins for forming a foamed resin layer of an expansion-molded article shown in Table 2 and their corresponding resins for forming a resin layer of an expansion-molded article shown in Table 3 were separately used to attempt to product a multi-layer parison in the same manner as in Example 1. However, the foamed resin layer was shrunk due to cell collapse and scarcely expanded, and so any good multi-layer parison was not obtained. This multi-layer parison was molded in the same manner as in Example 1. However, no expansion-molded article was obtained.

As described above, the expansion-molded articles according to the present invention are such that the thickness and density of the foamed polypropylene resin layer making up a multi-layer expansion-molded article of a polypropylene resin can be optionally selected or

controlled by adopting the above-described constitution,  
and so those having a foamed resin layer having a great  
thickness and a high expansion ratio may also be  
provided. In addition, the expansion-molded articles  
5 according to the present invention are uniform in the  
thickness of the polypropylene resin layer making up the  
molded articles and good in appearance without causing a  
scale pattern, cracks, holes and/or the like in the  
surface of the resin layer.

10           The production process of an expansion-molded  
article according to the present invention comprises  
molding a multi-layer parison having a resin layer on  
the outer surface of a foamed resin layer in a mold in  
such a manner that at least part of the opposed  
15 innermost foamed layers of the multi-layer parison are  
fusion-bonded to each other. Therefore, complicated  
production processes and apparatus like the conventional  
process for obtaining a multi-layer expansion-molded  
article of a polypropylene resin of the structure that a  
20 foamed resin layer has a skin layer on its surface are  
unnecessary. In addition, specific polypropylene resins  
are used as base resins, thereby improving the extension  
and drawdown tendency of the resin layer of the multi-  
layer parison and moreover improving the expansion and  
25 drawdown tendency of the foamed resin layer. Therefore,  
multi-layer expansion-molded articles of a polypropylene  
resin that have heretofore been difficult to obtain by

The expansion-molded articles according to the present invention are excellent in lightweight property, heat resistance, chemical resistance, recycling ability, heat insulating property, cushioning property, damping property, mechanical strength such as compressive, tensile and bending strength. The expansion-molded articles according to the present invention are also suitable for use as deep-draw expansion-molded articles and large-sized expansion-molded articles, and can be utilized as, to say nothing of returnable containers, light-weight heat insulating panels for floor, door or the like; structural members for automobile (particularly, shock-absorbing materials for automobile), such as pillars, bumpers, instrument panels, spoilers, fenders, side steps, door trims, grille guards and trunk boards; desks; chairs; floats; surfboards; pallets; etc.



## WHAT IS CLAIMED IS:

1. A multi-layer expansion-molded article of a polypropylene resin, which is obtained by molding a multi-layer parison comprising a foamed resin layer and a resin layer provided on the outer side of the foamed resin layer in a mold in such manner that at least part of the opposed inner surfaces of foamed resin layer in the parison are fusion-bonded to each other, and has a polypropylene resin layer on the surface of a foamed polypropylene resin layer, wherein a melt tension, MT (gf) and a melt flow rate, MFR (g/10 min) obtained by measurement to a polypropylene resin forming the foamed resin layer in the expansion-molded article satisfy the following relationship (1), and a melt tension, MT (gf) and a melt flow rate, MFR (g/10 min) obtained by measurement to a polypropylene resin forming the resin layer on the surface of the foamed resin layer satisfy the following relationship (2) when the melt flow rate, MFR is at least 0.3 (g/10 min), or the melt tension, MT is at least 10 (gf) when the melt flow rate, MFR is not lower than 0.2 (g/10 min), but lower than 0.3 (g/10 min):

$$\log MT > -0.74 \log MFR + 0.66 \quad (1)$$

$$\log MT > -1.02 \log MFR + 0.47 \quad (2)$$

2. A multi-layer expansion-molded article of a

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polypropylene resin, which is obtained by molding a multi-layer parison comprising a foamed resin layer and a resin layer provided on the outer side of the foamed resin layer in a mold in such manner that at least part of the opposed inner surfaces of foamed resin layer in the parison are fusion-bonded to each other, and has a polypropylene resin layer on the surface of a foamed polypropylene resin layer, wherein a melt tension, MT (gf) and a melt flow rate, MFR (g/10 min) obtained by measurement to a polypropylene resin forming the foamed resin layer in the expansion-molded article satisfy the following relationship (3), and a melt tension, MT (gf) and a melt flow rate, MFR (g/10 min) obtained by measurement to a polypropylene resin forming the resin layer on the surface of the foamed resin layer satisfy the following relationship (4) when the melt flow rate, MFR is at least 0.5 (g/10 min), or the melt tension, MT is at least 10 (gf) when the melt flow rate, MFR is not lower than 0.2 (g/10 min), but lower than 0.5 (g/10 min):

$$\log MT > -0.74 \log MFR + 0.79 \quad (3)$$

$$\log MT > -1.02 \log MFR + 0.69 \quad (4)$$

3. The multi-layer expansion-molded article of the polypropylene resin according to Claim 1 or 2, wherein the thickness of the resin layer formed on the surface of the foamed resin layer is 100  $\mu\text{m}$  to 10 mm, and the

overall density of the expansion-molded article is 20 to 400 kg/m<sup>3</sup>.

4. The multi-layer expansion-molded article of the  
5 polypropylene resin according to Claim 1 or 2, wherein  
the area ratio of the fusion-bonded portion in the inner  
surface of the expansion-molded article is at least 25%.

5. The multi-layer expansion-molded article of the  
10 polypropylene resin according to Claim 1 or 2, wherein  
the area ratio of the fusion-bonded portion in the inner  
surface of the expansion-molded article is at least 60%.

6. The multi-layer expansion-molded article of the  
15 polypropylene resin according to Claim 1 or 2, wherein  
the area ratio of the fusion-bonded portion in the inner  
surface of the expansion-molded article is at least 80%.

7. The multi-layer expansion-molded article of the  
20 polypropylene resin according to Claim 1 or 2, wherein  
the area ratio of the fusion-bonded portion in the inner  
surface of the expansion-molded article is at least 95%.

8. The multi-layer expansion-molded article of the  
25 polypropylene resin according to Claim 1 or 2, which  
further has a skin layer formed of a synthetic resin on  
the outer side of the resin layer.

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9. A process for producing a multi-layer expansion-molded article of a polypropylene resin, which comprises co-extruding a foamable molten resin obtained  
 5 by adding a foaming agent to a polypropylene resin whose melt tension, MT (gf) and melt flow rate, MFR (g/10 min) satisfy the following relationship (1), and melting and kneading the resultant mixture, and a molten resin obtained by melting and kneading a polypropylene resin  
 10 whose melt flow rate, MFR (g/10 min) is at least 0.3 (g/10 min), and whose melt tension, MT (gf) and melt flow rate, MFR (g/10 min) satisfy the following relationship (2), or whose melt flow rate, MFR is not lower than 0.1 (g/10 min), but lower than 0.3 (g/10 min)  
 15 and whose melt tension, MT is at least 10 (gf), thereby forming a multi-layer parison having a resin layer composed of the molten resin on the surface of a foamed resin layer obtained by expanding the foamable molten resin, and then molding the multi-layer parison in a  
 20 mold in such manner that at least part of the opposed inner surfaces of foamed resin layer in the parison are fusion-bonded to each other to obtain an expansion-molded article having a polypropylene resin layer on the surface of a foamed polypropylene resin layer:

$$25 \quad \log MT > -0.74 \log MFR + 0.66 \quad (1)$$

$$\log MT > -1.02 \log MFR + 0.47 \quad (2)$$

10. A process for producing a multi-layer expansion-molded article of a polypropylene resin, which comprises co-extruding a foamable molten resin obtained by adding a foaming agent to a polypropylene resin whose melt tension, MT (gf) and melt flow rate, MFR (g/10 min) satisfy the following relationship (5), and melting and kneading the resultant mixture, and a molten resin obtained by melting and kneading a polypropylene resin whose melt flow rate, MFR (g/10 min) is at least 0.8 (g/10 min), and whose melt tension, MT (gf) and melt flow rate, MFR (g/10 min) satisfy the following relationship (6), or whose melt flow rate, MFR is not lower than 0.2 (g/10 min), but lower than 0.8 (g/10 min) and whose melt tension, MT is at least 10 (gf), thereby forming a multi-layer parison having a resin layer composed of the molten resin on the surface of a foamed resin layer obtained by expanding the foamable molten resin, and then molding the multi-layer parison in a mold in such manner that at least part of the opposed inner surfaces of foamed resin layer in the parison are fusion-bonded to each other to obtain an expansion-molded article having a polypropylene resin layer on the surface of a foamed polypropylene resin layer:

$$\log MT > -0.74 \log MFR + 1.14 \quad (5)$$

$$\log MT > -1.02 \log MFR + 0.90 \quad (6)$$

11. The production process according to Claim 9 or

10, wherein the polypropylene resin used for the foamable molten resin has an MFR of 0.3 to 20 g/10 min.

12. A container formed of the multi-layer expansion-molded article of the polypropylene resin according to Claim 1 or 2, wherein the overall density of the container is 30 to 400 kg/m<sup>3</sup>, and the thickness of the polypropylene resin layer is 200 μm to 5 mm.
13. A shock-absorbing material for automobile formed of the multi-layer expansion-molded article of the polypropylene resin according to Claim 1 or 2, wherein the overall density of the shock-absorbing material is 25 to 300 kg/m<sup>3</sup>, and the thickness of the polypropylene resin layer is 200 μm to 7 mm.
14. A member for automobile formed of the multi-layer expansion-molded article of the polypropylene resin according to Claim 1 or 2.
15. The member for automobile according to Claim 14, wherein the member for automobile is a member selected from among a bumper, pillar, instrument panel, spoiler, fender, side step, door trim, grille guard and trunk board.

## ABSTRACT OF THE DISCLOSURE

Disclosed herein is a multi-layer expansion-molded article of a polypropylene resin, which is obtained by molding a multi-layer parison comprising a foamed resin layer and a resin layer provided on the outer side of the foamed resin layer in a mold in such manner that at least part of the opposed inner surfaces of foamed resin layer in the parison are fusion-bonded to each other, and has a polypropylene resin layer on the surface of a foamed polypropylene resin layer, wherein a melt tension, MT (gf) and a melt flow rate, MFR (g/10 min) obtained by measurement to a polypropylene resin forming the foamed resin layer in the expansion-molded article satisfy the following relationship (1), and a melt tension, MT (gf) and a melt flow rate, MFR (g/10 min) obtained by measurement to a polypropylene resin forming the resin layer on the surface of the foamed resin layer satisfy the following relationship (2) when the melt flow rate, MFR is at least 0.3 (g/10 min), or the melt tension, MT is at least 10 (gf) when the melt flow rate, MFR is not lower than 0.2 (g/10 min), but lower than 0.3 (g/10 min):

$$\log MT > -0.74 \log MFR + 0.66 \quad (1)$$

$$\log MT > -1.02 \log MFR + 0.47 \quad (2)$$

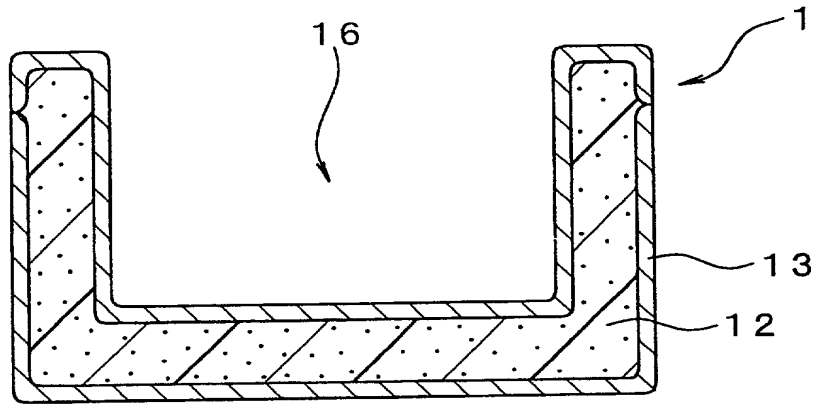


FIG. 1A

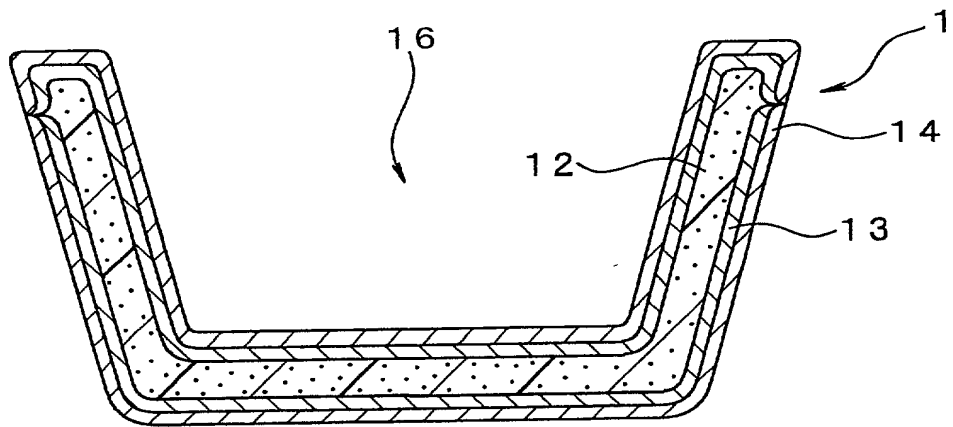


FIG. 1B



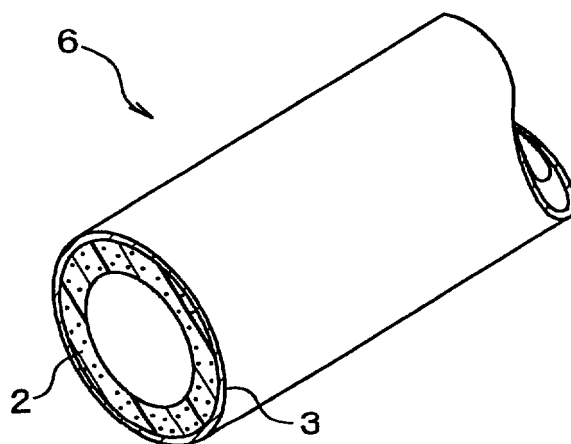


FIG. 2A

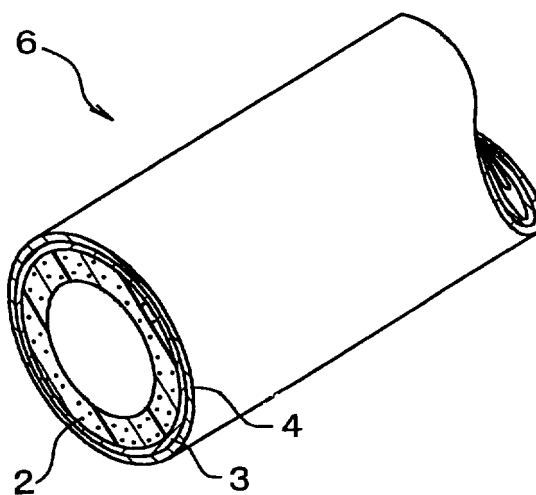


FIG. 2B

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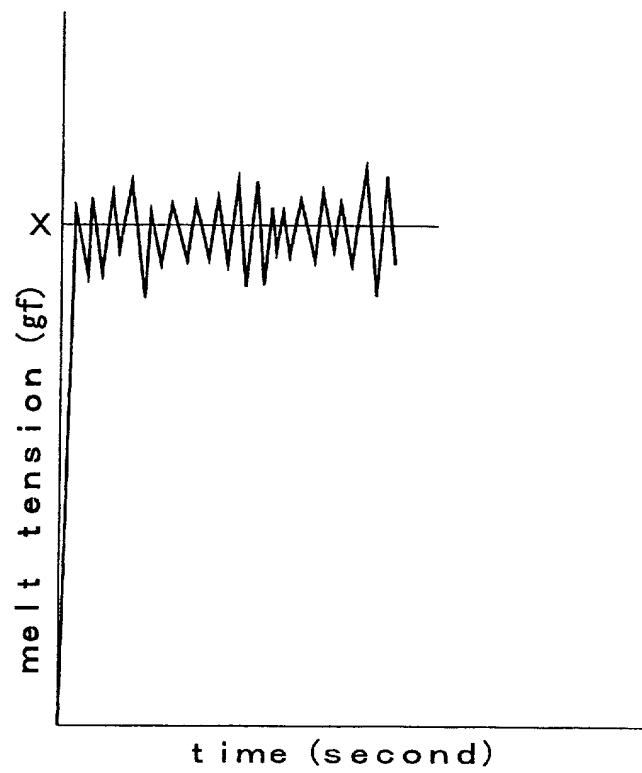


FIG. 3

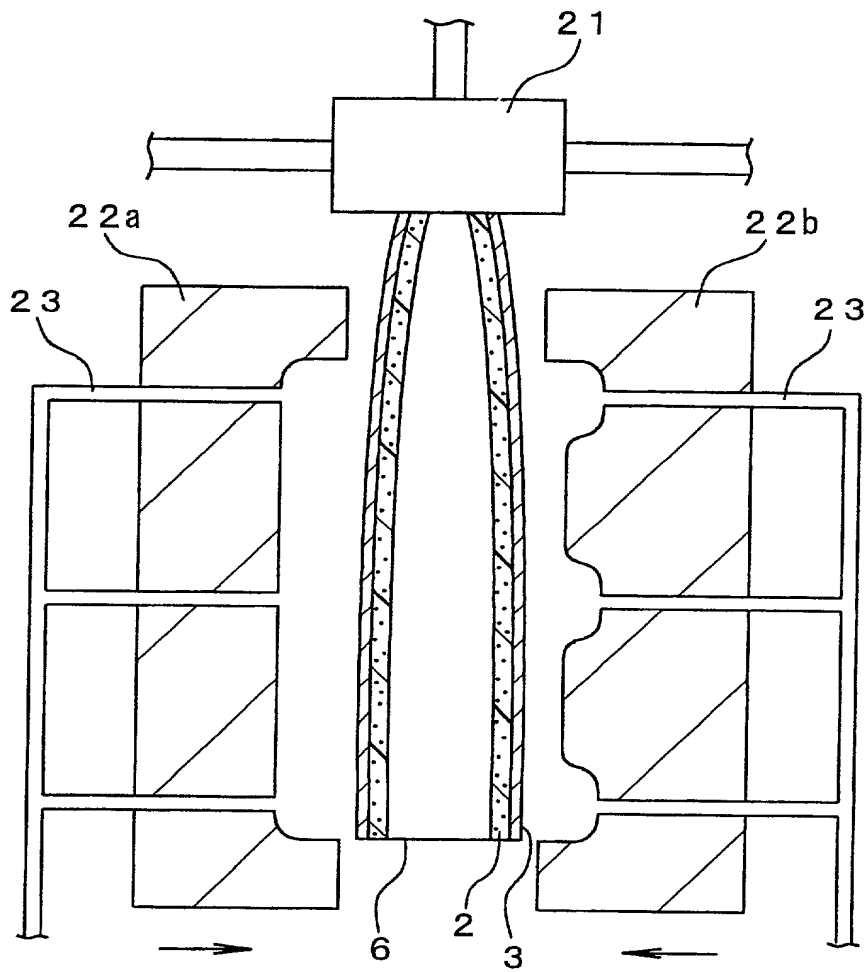


FIG. 4

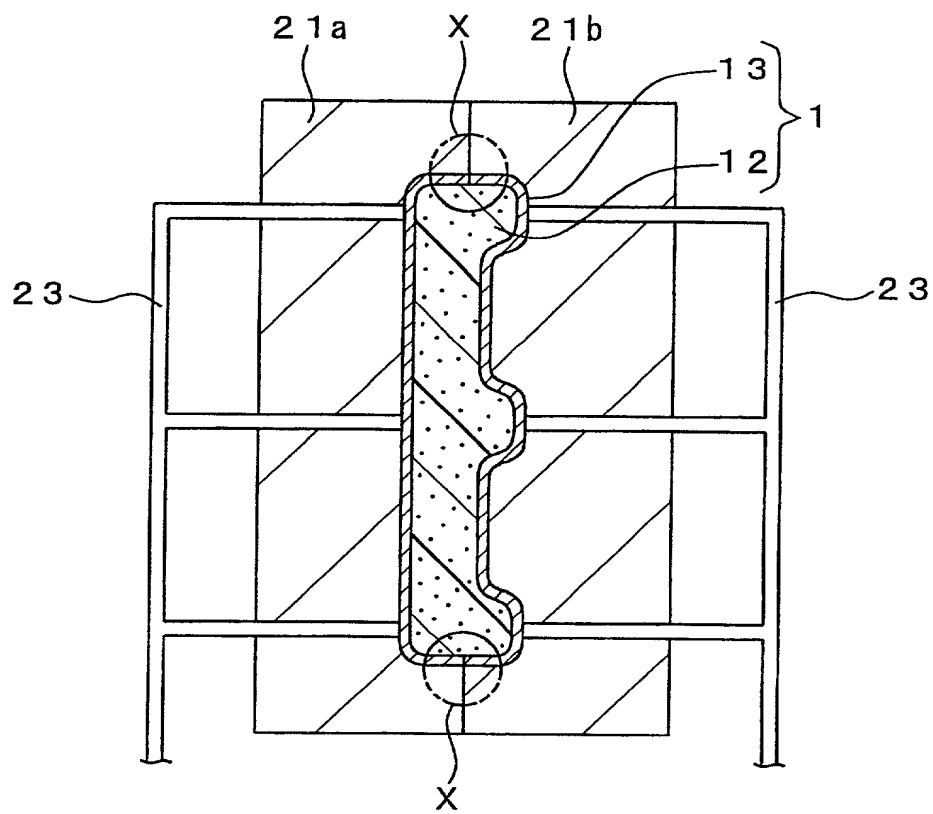


FIG. 5

FIG. 6B

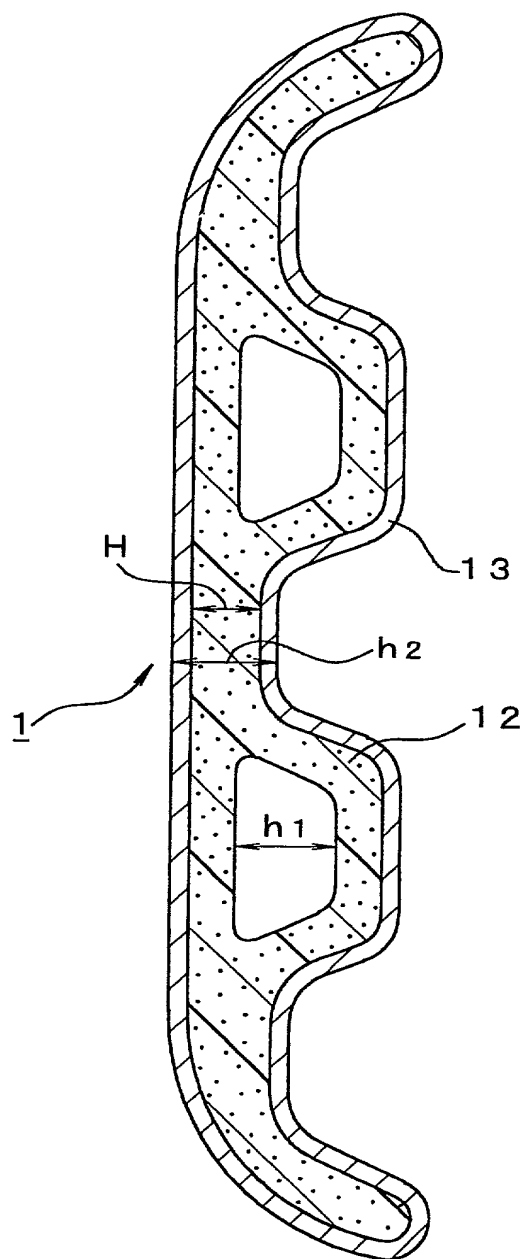


FIG. 7

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## DECLARATION AND POWER OF ATTORNEY FOR PATENT APPLICATION

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled MULTI-LAYER EXPANSION-MOLDED ARTICLE OF POLYPROPYLENE RESIN, PRODUCTION PROCESS THEREOF AND CONTAINER, SHOCK-ABSORBING MATERIAL FOR AUTOMOBILE AND AUTOMOTIVE MEMBER FORMED OF THE MULTI-LAYER EXPANSION-MOLDED ARTICLE OF POLYPROPYLENE RESIN, the specification of which:

(check one) ☒ is attached hereto ☐ was filed on \_\_\_\_\_ as Application Serial No. \_\_\_\_\_ and was amended on \_\_\_\_\_ (if applicable).

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the examination of this application in accordance with Title 37, Code of Federal Regulations, § 1.56(a).

I hereby claim foreign priority benefits under Title 35, United States Code § 119 or any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed:

## Prior Foreign Application(s)

Prior Foreign Application(s)			Priority claimed	
<u>225346/1999</u>	<u>Japan</u>	<u>09 August 1999</u>	<input checked="" type="checkbox"/>	<input type="checkbox"/>
(NUMBER)	(COUNTRY)	(D/M/Y)	YES	NO
<u>                    </u>	<u>                    </u>	<u>                    </u>	<input type="checkbox"/>	<input type="checkbox"/>
(NUMBER)	(COUNTRY)	(D/M/Y)	YES	NO
<u>                    </u>	<u>                    </u>	<u>                    </u>	<input type="checkbox"/>	<input type="checkbox"/>
(NUMBER)	(COUNTRY)	(D/M/Y)	YES	NO

I hereby claim the benefit under Title 35, United States Code, § 120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, § 112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, § 156(a) which occurred between the filing date of the prior application and the national or PCT international filing date of this applications:

<u>                                    </u>	<u>                                    </u>	<u>                                    </u>
(APPLICATION SERIAL NO.)	(FILING DATE)	(STATUS)
		(PATENTED, PENDING, ABANDONED)

<u>                                    </u>	<u>                                    </u>	<u>                                    </u>
(APPLICATION SERIAL NO.)	(FILING DATE)	(STATUS)
		(PATENTED, PENDING, ABANDONED)

POWER OF ATTORNEY: As a named inventor, I hereby appoint the following attorneys and/or agents to prosecute this application and transact all business in the Patent and Trademark Office connected therewith:

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I further declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine and imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or document or any patent issuing thereon.

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[X] ADDITIONAL INVENTORS ARE BEING NAMED ON SEPARATELY NUMBERED SHEETS ATTACHED HERETO

OFFICE OF THE SECRETARY OF COMMERCE



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